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Table 3. Clinical Laboratory Director Reporting Requirements

①	Arboviruses	☎ , *	Haemophilus influenzae, type b, isolated from a normally sterile site	=	Plasmodium spp.
≜, ≊ , *	Bacillus anthracis	᠍,₩	<i>Haemophilus influenzae</i> , other, isolated from a normally sterile site	=	Respiratory synctial virus
☎ , *	Bordetella pertussis	=	Hantavirus	* ,+	Rubella virus and anti-rubella-IgM serologies
① ,*	Brucella spp.	= 1	Hepatitis A virus (anti-HAV-IgM serologies)	① ,*	Salmonella spp.
) ,*	Burkholderia mallei and B. pseudomalle	ri=1	Hepatitis B virus (anti-Hepatitis B core-IgM serologies, Hepatitis B surface or envelope antigen serologies, or detection of viral nucleic acid)	~	SARS-associated corona virus
="	Campylobacter spp.	<u>=</u> 1	Hepatitis C virus) ,*	Shigella spp.
≡.	CD ₄ -T-lymphocyte count of fewer than 200 per microliter of whole blood or CD ₄ -T-lymphocyte percentage of total lymphocytes of less than 14%	<u>=</u> 1	Hepatitis D virus	=	Streptococcus Group A, isolated from a normally sterile site
=	Chlamydia trachomatis	<u> </u>	Hepatitis E virus (anti-HEV-IgM serologies)	=	Streptococcus Group B, isolated from a normally sterile site in an infant younger than 90 days of age
⊕, જ	Clostridium botulinum toxin (botulism)	=	HIV (by culture, antigen, antibodies to the virus, or detection of viral nucleic acid)	≡ , *	Streptococcus pneumoniae and its drug sensitivity pattern, isolated from a normally sterile site
=	Coccidioides spp., by culture or serologies	==	HIV—any test result for an infant (by culture, antigen, antibodies to the virus, or detection oviral nucleic acid)		Treponema pallidum (syphilis)
1	Coxiella burnetti	=	Influenza virus	=	Trypanosoma cruzi (Chagas disease)
=	Cryptosporidium spp.	≡,*	Legionella spp. (culture or DFA)	① ,*	Vancomycin-resistant or Vancomycin- intermediate <i>Staphylococcus aureus</i>
①	Cyclospora spp.	① ,*	<i>Listeria</i> spp., isolated from a normally sterile site	① ,*	Vancomycin resistant Staphylococcus epidermidis
=	Dengue virus	* ,+	Measles virus and anti-measles-IgM serologies	⊜,☎	Variola virus (smallpox)
≜,☎	Emerging or exotic disease agent	2	Methicillin-resistant <i>Staphylococcus aureus</i> , isolated from a normally sterile site	① , *	Vibrio spp.
="	Entamoeba histolytica	1 ,+	Mumps virus and anti-mumps-IgM serologies	≙,☎	Viral hemorrhagic fever agent
①	Escherichia coli O157:H7	= , * ³	Mycobacterium tuberculosis complex and its drug sensitivity pattern	=	West Nile virus
1 ,*	Escherichia coli, Shiga-toxin producing	=	Neisseria gonorrhoeae) ,*	Yersinia spp. (other than Y. pestis)
⊕, ☎ , *	Francisella tularensis	2 ,*	Neisseria meningitidis, isolated from a normally sterile site	⊕,≊,*	Yersinia pestis (plague)
		=	Norovirus		
From I	December 1, 2008 until February 28, 2010	<u>=14</u>	Clostridium difficile toxin		

Key:

- Submit a report immediately after receiving one specimen for detection of the agent. Report receipt of subsequent specimens within five working days after receipt.
- Submit a report within 24 hours after obtaining a positive test result.
- Submit a report within one working day after obtaining a positive test result.
- Submit a report within five working days after obtaining a positive test result or a test result specified in Table 3.
- * Submit an isolate of the organism for each positive culture to the Arizona State Laboratory at least once each week, as applicable.
- + For each positive test result, submit a specimen to the Arizona State Laboratory within 24 hours after obtaining the positive test result.
- When reporting a positive result for any of the specified tests, report the results of all other tests performed for the subject as part of the disease panel.
- Submit a report only when an initial positive result is obtained for an individual.
- Submit an isolate of the organism only when an initial positive result is obtained for an individual, when a change in resistance pattern is detected, or when a positive result is obtained ≥ 12 months after the initial positive result is obtained for an individual.
- For every eighth positive test result, submit a specimen to the Arizona State Laboratory or the Department's designee. A clinical laboratory director may aggregate specimens and submit specimens every five working days, rather than submitting individual specimens.